

Express Mail No. EV449562872US PATENT

## IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicants

Jeffery R. Raymond et al.

Application No.

10/624,946

Confirmation No.

7685

Filed

July 21, 2003

For

3-NITROGEN-6,7-DIOXYGEN STEROIDS AND USES RELATED

**THERETO** 

Art Unit

1614

Docket No.

480117.407C1

Date

September 2, 2004

Mail Stop PG PUB Commissioner for Patents P.O. Box 1450 Alexandria, VA 22313-1450

## REQUEST FOR CORRECTED APPLICATION PUBLICATION

## Commissioner for Patents:

In response to the Notice of Publication of Application, dated July 15, 2004, Applicants respectfully request correction and republication of the above-identified application which they believe to contain material errors.

In particular, lines 1-2 of Claim 38 in the published patent application **incorrectly** read as follows:

38. A compound of claim 1 wherein  $-N(R I)(R^2)$  is in a salt form and the salt is a halogen or acetate salt.

U.S. Application No. 10/744,857432

These lines should **correctly** read as follows:

38. A compound of claim 1 wherein  $-N(R^1)(R^2)$  is in a salt form and the salt is a halogen or acetate salt.

To assist the Office in identifying the error, enclosed is a copy of page 52 of the published patent application, with a red circle around the language in the claim that was incorrectly printed. A copy of the corresponding page of the patent application (Page 124) as originally filed is likewise attached, with a red circle around the correct language in the claim.

This error is material in that it affects the scope of Claim 38.

Applicants respectfully request that the above-noted material error be corrected and that the application be republished by the Office. As the error was made on the part of the USPTO, no fee should be due.

The Office is requested to contact the undersigned at (206) 622-4900 should any additional issue require attention prior to a final decision being made by the Office.

Respectfully submitted,

Seed Intellectual Property Law Group PLLC

Carol J. Roth

Registration No. 32,783

CJR:cw Enclosures:

Transmittal Form
Page 124 of the Application as filed
Page 52 of the Published Application

701 Fifth Avenue, Suite 6300 Seattle, Washington 98104-7092

Phone: (206) 622-4900 Fax: (206) 682-6031

510285

TRAN	SMITTAL ORM r all correspondence initial filing)	Application Number Filing Date First Named Inventor Group Art Unit	nformation unless it displays a valid OMB control number 10/624,946  July 21, 2003  Jeffery R. Raymond
(To be used for after	ORM r all correspondence	First Named Inventor Group Art Unit	
(To be used fo	r all correspondence	Group Art Unit	Jeffery R. Raymond
after		<del> </del>	
after		Evaminar Nama	1614
Fee Transmit		Lxammer mame	
Fee Transmit		Attorney Docket No.	480117.407C1
Fee Transmit	ENC	CLOSURES (check all that ap	ply)
Extension of Express Abar Request Information D Statement; For Cited Referent Certified Copy Document(s) Response to Under 37 C.F.	Althed Response II Ideclaration(s) Fime Request Idenment	Drawing(s) Request for Corrected Filing Receipt Licensing-related Papers Petition Petition to Convert to a Provisional Application Power of Attorney, Revocation, Change of Correspondence Address Declaration Statement under 37 CFR 3.73(b) Terminal Disclaimer Request for Refund	CD(s), Number of CD(s)  After Allowance Communication to Group Appeal Communication to Board of Appeals and Interferences Appeal Communication to Group (Appeal Notice, Brie Reply Brief) Proprietary Information Status Letter Return Receipt Postcard Additional Enclosure(s) (please identify below): Request for Corrected Application Publication: Copy of Page 52 of the Application as filed; Copy of Page 124 of the Published Application
Remarks			
	SIGNATURE	OF APPLICANT, ATTORNEY	, OR AGENT
Individual Name	Carol J. Roth		Customer Number
Registration		. 50,922	00500
Signature Carol		Roth	
Date	September 2, 2		
		ICATE OF TRANSMISSION/M	
with the United S	ates Postal Servic	ce with sufficient postage as fir	itted to the USPTO or deposited st class mail in an envelope dria, VA 22313-1450 on the date
SHOWIT DEIGW.			
Typed or printed	name ~~ <b>S</b> @	ent via Express Mail ~~	

19. A compound of claim 1 wherein 17 is substituted with  $-OR^6$  or -O, wherein  $R^6$  is hydrogen.

20. A compound of claim 1 wherein R<sup>1</sup> is selected from  $-C(=O)-R^7$ ,  $-C(=O)NH-R^7$ ;  $-SO_2-R^7$ ; wherein R<sup>1</sup> is selected from alkyl, heteroalkyl, aryl and heteroaryl.

21. A compound of claim 20 wherein  $R^7$  is selected from  $C_{1-10}$ hydrocarbyl.

22. A compound of claim 20 wherein R<sup>7</sup> comprises biotin.

23. A compound of claim 1 wherein  $(R^1)(R^2)N$ — is selected from

24. A compound of claim 1 wherein  $R^1$  is hydrogen and  $R^2$  comprises a carbocycle.

25. A compound of claim 24 wherein the carbocycle is phenyl.

26. A compound of claim 25 wherein R<sup>2</sup> is selected from 3-methylphenyl; 4-hydroxyphenyl; and 4-sulfonamidephenyl.

27. A compound of claim 1 wherein  $R^1$  is hydrogen and  $R^2$  comprises a  $C_{1-10}$ hydrocarbyl.

28. A compound of claim 1 wherein  $R^1$  is hydrogen and  $R^2$  is heteroalkyl.

29. A compound of claim 28 wherein  $R^2$  is selected from  $C_{1-10}$ alkyl-W— $C_{1-10}$ alkylene- wherein W is selected from O and NH; HO— $C_{1-10}$ alkylene-; and HO— $C_{1-10}$ alkylene-W— $C_{1-10}$ alkylene- where W is selected from O and NH.

30. A compound of claim 1 wherein  $R^1$  is hydrogen and  $R^2$  is —CH<sub>2</sub>— $R^7$  wherein  $R^7$  is selected from alkyl, heteroalkyl, aryl and heteroaryl.

31. A compound of claim 30 wherein R<sup>7</sup> is selected from alkyl-substituted phenyl; halogen-substituted phenyl; alkoxy-substituted phenyl; aryloxy-substituted phenyl; and nitro-substituted phenyl.

32. A compound of claim 1 wherein each of R<sup>1</sup> and R<sup>2</sup> is hydrogen.

33. A compound of claims 1 or 32 wherein each of  $\mathbb{R}^3$  and  $\mathbb{R}^4$  is hydrogen.

34. A compound of claims 32 or 33 where the carbon at numeral 17 is substituted with

(a) one of the following:  $C(R^{5a})(R^{5a})$ ,  $=C=C(R^{5a})(R^{5a})$ , and  $-C(R^{5a})(R^{5a})(C(R^{5a})(R^{5a}))_n$ — wherein n ranges from 1 to about 6; or

(b) two of the following, which are independently selected: —X, —N(R¹)(R²), and —R⁵a;

where R<sup>5a</sup> at each occurrence is independently selected from H, X, and C<sub>1-30</sub> organic moiety that may optionally contain at least one heteroatom selected from the group consisting of boron, halogen, nitrogen, silicon and sulfur; where two geminal R<sup>5</sup> groups may together form a ring with the carbon atom to which they are both bonded.

35. A compound of claim 1 wherein  $R^3$  and  $R^4$  together form a ketal of the structure

36. A compound of claim 1 wherein —OR3 and —OR4 have the stereochemistry shown

37. A compound of claim 1 wherein  $-N(R^1)(R^2)$  is in a

38. A compound of claim 1 wherein —N(R I)(R<sup>2</sup>) is in a salt form and the salt is a halogen or acetate salt.

39. A compound of claim 1 which is a prograg of the formula shown in claim 1.

40. A compound of claim 1 and pharmaceutically acceptable salts, solvates, stereoisomers but not prodrugs thereof, in isolation or in mixture.

41. A compound of claim 1 wherein at least one of the carbons at numerals 10 and 13 are substituted with methyl.

42. A compound of claim 1 wherein each of R<sup>1</sup> and R<sup>2</sup> are independently selected from hydrogen and organic groups having 1-20 carbons and optionally containing 1-5 heteroatoms selected from nitrogen, oxygen, silicon, and sulfur.

43. A compound of claim 1 wherein

R¹ and R² are independently selected from hydrogen, R<sup>8</sup>, R<sup>9</sup>, R<sup>10</sup>, R<sup>11</sup> and R<sup>12</sup> where R<sup>8</sup> is selected from alkyl, heteroalkyl, aryl and heteroaryl; R° is selected from (R<sup>8</sup>)<sub>r</sub>-alkylene, (R<sup>8</sup>)<sub>r</sub>-heteroalkylene, (R<sup>8</sup>)<sub>r</sub>-arylene and (R<sup>8</sup>)<sub>r</sub>-heteroarylene; R<sup>10</sup> is selected from (R°)<sub>r</sub>-alkylene, (R°)<sub>r</sub>-heteroalkylene, (R°)<sub>r</sub>-arylene, and (R°)<sub>r</sub>-heteroarylene; R<sup>11</sup> is selected from (R<sup>10</sup>)<sub>r</sub>-alkylene, (R<sup>10</sup>)<sub>r</sub>-heteroarylene, R<sup>12</sup> is selected from (R<sup>11</sup>)<sub>r</sub>-alkylene, heteroalkylene, (R<sup>11</sup>)<sub>r</sub>-arylene, and (R<sup>11</sup>)<sub>r</sub>-heteroarylene, and r is selected from 0, 1, 2, 3, 4 and 5, with the proviso that R¹ and R² may join to a common atom so as to form a ring with the common atom.

35. A compound of claim 1 wherein R<sup>3</sup> and R<sup>4</sup> together form a ketal of the structure

36. A compound of claim 1 wherein -OR<sup>3</sup> and -OR<sup>4</sup> have the stereochemistry shown

- 37. A compound of claim 1 wherein  $-N(R^1)(R^2)$  is in a salt form.
- 38. A compound of claim 1 wherein  $-N(R^1)(R^2)$  is in a salt form and the salt is a halogen or acetate salt.
  - 39. A compound of claim 1 which is a prodrug of the formula shown in claim
- 40. A compound of claim 1 and pharmaceutically acceptable salts, solvates, stereoisomers but not prodrugs thereof, in isolation or in mixture.

1.